

| | |
|---------------|--|
| Title | Entropic stabilization of the tryptophan synthase α -subunit from a hyperthermophile, <i>Pyrococcus furiosus</i> : X-ray analysis and calorimetry |
| Author(s) | Yamagata, Yuriko; Ogasahara, Kyoko; Hioki, Yusaku et al. |
| Citation | Journal of Biological Chemistry. 276(14) p.11062-p.11071 |
| Issue Date | 2001-04 |
| oaire:version | VoR |
| URL | https://hdl.handle.net/11094/73647 |
| rights | © the American Society for Biochemistry and Molecular Biology. |
| Note | |

Osaka University Knowledge Archive : OUKA

<https://ir.library.osaka-u.ac.jp/>

Osaka University

Entropic Stabilization of the Tryptophan Synthase α -Subunit from a Hyperthermophile, *Pyrococcus furiosus*

X-RAY ANALYSIS AND CALORIMETRY*

Received for publication, November 2, 2000, and in revised form, December 1, 2000
Published, JBC Papers in Press, December 14, 2000, DOI 10.1074/jbc.M009987200

Yuriko Yamagata‡, Kyoko Ogasahara§, Yusaku Hioki§, Soo Jae Lee§, Atsushi Nakagawa§, Haruki Nakamura§, Masami Ishida¶, Seiki Kuramitsu||, and Katsuhide Yutani§**

From the ‡Graduate School of Pharmaceutical Sciences, and §Institute for Protein Research, Osaka University, Yamadaoka, Suita, Osaka 565-0871, Japan, the ¶Tokyo University of Fisheries, Konan Minato-ku, Tokyo 108-8477, Japan, and the ||Graduate School of Sciences, Osaka University, Machikaneyama, Toyonaka, Osaka 560-0043, Japan

The structure of the tryptophan synthase α -subunit from *Pyrococcus furiosus* was determined by x-ray analysis at 2.0-Å resolution, and its stability was examined by differential scanning calorimetry. Although the structure of the tryptophan synthase $\alpha_2\beta_2$ complex from *Salmonella typhimurium* has been already determined, this is the first report of the structure of the α -subunit alone. The α -subunit from *P. furiosus* (*Pf*- α -subunit) lacked 12 and 6 residues at the N and C termini, respectively, and one residue each in two loop regions as compared with that from *S. typhimurium* (*St*- α -subunit), resulting in the absence of an N-terminal helix and the shortening of a C-terminal helix. The structure of the *Pf*- α -subunit was essentially similar to that of the *St*- α -subunit in the $\alpha_2\beta_2$ complex. The differences between both structures were discussed in connection with the higher stability of the *Pf*- α -subunit and the complex formation of the α - and β -subunits. Calorimetric results indicated that the *Pf*- α -subunit has extremely high thermostability and that its higher stability is caused by an entropic effect. On the basis of structural information of both proteins, we analyzed the contributions of each stabilization factor and could conclude that hydrophobic interactions in the protein interior do not contribute to the higher stability of the *Pf*- α -subunit. Rather, the increase in ion pairs, decrease in cavity volume, and entropic effects due to shortening of the polypeptide chain play important roles in extremely high stability in *Pf*- α -subunit.

Prokaryotic tryptophan synthase, which catalyzes the last processes in the biosynthesis of tryptophan, is a multienzyme $\alpha_2\beta_2$ complex composed of nonidentical α - and β -subunits. The separate α - and β_2 -subunits catalyze inherent reactions termed α and β reactions, respectively. When the α - and β_2 -subunits combine to form the $\alpha_2\beta_2$ complex, the enzymatic activity of each subunit is stimulated by 1 to 2 orders of magnitude (1). The $\alpha_2\beta_2$ complex has been studied as an excellent model

system for seeking answers to important questions in protein-protein interaction, especially in multifunctional enzymes. In 1988 (2) the three-dimensional structure of the tryptophan synthase $\alpha_2\beta_2$ complex from *Salmonella typhimurium* was determined by x-ray analysis. However, the structure of the α - or β_2 -subunit alone has not yet been determined. To elucidate the molecular basis of the mutual activation of the subunit interaction due to the formation of the $\alpha_2\beta_2$ complex, we need to know the structures of the α - or β_2 -subunits alone as well as that of the complex. Although the crystallization of each subunit from *S. typhimurium* and *Escherichia coli* has been tried for many years (3), the report of the x-ray structure has not yet appeared. Recently, the structures of a number of proteins from hyperthermophiles have been successfully determined by x-ray analysis. This seems due to the facts that proteins from hyperthermophiles are unusually stable and more easily form better crystals. Therefore, the *trpA* and *trpB* genes, coding for tryptophan synthase, were cloned from *Pyrococcus furiosus* and their products were overexpressed in *E. coli*. The purified α -subunit from *P. furiosus* easily grew a suitable crystal for x-ray analysis, and the crystal structure could be determined at a resolution of 2.0 Å as described in this paper.

On the other hand, a fundamental understanding of the conformational stability of proteins still remains elusive. Proteins from hyperthermophiles should retain their native structures under extreme conditions, whereas the homologous proteins from mesophiles completely denature. Therefore, comparative studies of extremely thermostable proteins with their homologs might help us understand the stabilization mechanism of a general globular protein. Recently, structures and physicochemical properties of proteins from hyperthermophiles have been extensively studied. In several hyperthermophile proteins, an increased number of ion pairs and ion pair networks have been observed (4–13), which have been explained as the intrinsic changes for protein stability in hyperthermophiles. In the case of the tryptophan synthase α -subunit from *P. furiosus*, many ion pairs were also found. However, the effect of a surface salt bridge on the stability remains controversial even today; some reports have shown little contribution of a surface salt bridge to stability (14–19), whereas others have shown a favorable contribution (20–24). There are also reports on some hyperthermophile proteins without additional ion pairs (25–28). The conformation of general globular proteins is marginally maintained by the combination of many positive (such as hydrophobic interaction and hydrogen bond) and negative (such as entropic effect and steric hindrance) factors for stabilization (29). The enhanced stability of hyperthermophile proteins might originate from several attractive

* This work was supported in part by a grant-in-aid for special project research from the Ministry of Education, Science, and Culture of Japan (to K. Y.) and by the Structural Biology Sakabe Project (to K. Y., A. N., and Y. Y.). The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

** To whom correspondence should be addressed: Institute for Protein Research, Osaka University, 3-2 Yamadaoka, Suita, Osaka 565-0871, Japan. Tel.: 81-6-6879-8615; Fax: 81-6-6879-8616; E-mail: yutani@protein.osaka-u.ac.jp.

forces (30). However, it still remains unclear what causes the dramatic stabilization of proteins from hyperthermophiles. Understanding the molecular origin for stabilization of hyperthermophile proteins provides valuable insights into the problems of protein stability, protein folding, and protein engineering.

In this paper, to elucidate the stabilization mechanism of a protein from a hyperthermophile, the stability of the α -subunit from *P. furiosus* (*Pf*- α -subunit)¹ was examined by differential scanning calorimetry (DSC) and its structure by x-ray crystallography. The DSC data indicated that the higher stability of this protein was not caused by an enthalpic factor as compared with homologous mesophile proteins. The stabilization mechanism of the *Pf*- α -subunit is discussed on thermodynamic grounds based on structural information from x-ray analysis.

EXPERIMENTAL PROCEDURES

Purification of the α -Subunit from *P. furiosus*—The α -subunit of tryptophan synthase from *P. furiosus* (*Pf*- α -subunit) was overexpressed in *E. coli* strain JM109/p α 1974 containing only the *trpA* gene from *P. furiosus*.² The *E. coli* strain was routinely grown in 15 liters of LB medium supplemented with ampicillin of 100 mg/liter of culture medium. The production of the *Pf*- α -subunit was induced by addition of isopropyl- β -D(-)-thiogalactopyranoside at a concentration of 1 mM to the culture medium after 1-h incubation at 37 °C with shaking. Then the culture was continued for about 20 h at 37 °C with shaking.

Pelleted cells (wet weight of about 25 g) collected by centrifugation (5000 rpm for 10 min) were suspended in 100 ml of 20 mM potassium phosphate buffer (pH 7.0) containing 1 mM EDTA and 5 mM DTT. The cells were disrupted by sonication on ice for 5 min with cooling intervals for 10 min and repeating three times. The homogenized solution was heated in a water bath for 10 min at 75 °C (the temperature of the solution), under which the majority of the *E. coli* proteins precipitated, leaving the *Pf*- α -subunit in the soluble fraction. The resultant suspended solution was centrifuged at 15,000 rpm for 30 min at 4 °C to remove cell debris and denatured *E. coli* proteins. The supernatant solution was treated with ammonium sulfate of 40% saturation and centrifuged, and the precipitate was discarded. The *Pf*- α -subunit was precipitated by addition of ammonium sulfate at 90% saturation. The precipitate was dissolved in 50 ml of 25 mM potassium phosphate buffer (pH 7.0) with 1 mM EDTA and 5 mM DTT and dialyzed against the same buffer overnight at 4 °C.

The dialyzed sample was applied on a column (2.5 \times 20 cm) of DEAE-Sephacel (Amersham Pharmacia Biotech): the enzyme was eluted with a linear gradient (25–200 mM) of potassium phosphate buffer (pH 7.0) containing 1 mM EDTA and 5 mM DTT. Active fractions of the eluted solutions were concentrated using an Amicon PM10 membrane filter. Next, the protein was separated by gel filtration (Superdex TM200 26/60, Amersham Pharmacia Biotech). The active fractions collected were finally purified by ion exchange chromatography (SP-Sephacrose 26/16, Amersham Pharmacia Biotech) with a linear gradient of 25–100 mM phosphate buffer (pH 7.0) containing 1 mM EDTA and 5 mM DTT. The purified protein showed a single band on SDS-polyacrylamide gel electrophoresis.

Protein Concentration—The protein concentration was estimated from absorbance at 278.5 nm at pH 7.0, OD_{1%} = 6.92, using a cell with a light path length of 1 cm. The value was determined based on a protein assay by the Lowry method using bovine serum albumin as a standard protein.

Differential Scanning Calorimetry—DSC was carried out with a differential scanning calorimeter, MicroCal VP-DSC (Northampton) at a scan rate of 1 K/min. Prior to the measurements, the protein solution was dialyzed against the buffer used. The dialyzed sample was filtered through a 0.22- μ m pore size membrane and then degassed in vacuum. The buffers used were 20 mM glycine-HCl in the acidic region and 20 mM glycine-KOH in the alkaline region. The protein concentrations under measurement were 0.4–1.4 mg/ml. The DSC curves were analyzed by the Origin software from MicroCal (Northampton).

¹ The abbreviations used are: *Pf*- α -subunit, tryptophan synthase α -subunit from *P. furiosus*; *Ec*- α -subunit, tryptophan synthase α -subunit from *E. coli*; *St*- α -subunit, tryptophan synthase α -subunit from *S. typhimurium*; DSC, differential scanning calorimetry; DTT, dithiothreitol; MES, 4-morpholineethanesulfonic acid; ASA, accessible surface area.

² M. Ishida, T. Oshima, and K. Yutani, manuscript in preparation.

Crystallization and Data Collection—Crystals suitable for data collection were grown by the hanging drop vapor diffusion method at 10 °C from a reservoir solution containing 0.1 M MES-NaOH, pH 6.5, and 12% polyethylene glycol 20000. X-ray data from the native and mercury derivative crystals were collected at the beam line 18B of the Photon Factory, Tsukuba using a Weissenberg camera (32) at 293 K. The data set of the platinum derivative was collected on a Rigaku R-axis IV imaging plate using nickel-filtered double-mirror focused Cu-K α radiation from a Rigaku RU-200. The native and derivative data were processed and integrated by DENZO and scaled by SCALEPACK (33). The space group is C222₁, and the unit cell dimensions are $a = 73.013$, $b = 78.997$, and $c = 170.964$ Å (see Table II below). There were two molecules in the asymmetric unit with a volume per unit of molecular weight of the protein of 2.30 Å³/Da and a calculated solvent content of 53.4% (34).

Structure Determination and Refinement—Native, mercury, and platinum data sets were used for phase calculation by MIRAS (see Table II). The mercury position was identified in the difference Patterson map. Heavy atom sites in the platinum derivative were found in the difference Fourier maps calculated by using the phases from the first derivative with the program MLPHARE (35). The heavy atom parameters were refined, and the phases were calculated to 2.2 Å with the program SHARP (36). The initial MIR phases were improved by solvent flattening and noncrystallographic 2-fold symmetry averaging at the same resolution with the program SOLOMON (37). An initial model was built using the program O (38). Several cycles of rigid-body refinement, positional refinement, and simulated annealing were performed at 2.2-Å resolution with X-PLOR (39). The refinements were continued at 2.0 Å using CNS (40). Successive refinement with temperature factors and addition of solvents resulted in an R -value of 19.8% and an R_{free} of 24.9% for reflections in the resolution range 40–2.0 Å. R_{free} was calculated with 5% of the reflections. During refinements NCS restraints were enforced. The current model includes residues 1–166, 174–248 for molecule a and 1–169, 173–248 for molecule b, and 421 water molecules. Model geometry was analyzed with PROCHECK (41), and 96.4% of the nonglycine residues were in the most favorable region of the Ramachandran plot and 3.3% in the additionally allowed region. The final coordinates have been deposited in the Protein Data Bank (accession number 1GEQ).

Calculation of Electrostatic Interaction—The electrostatic free energy G_{el} of the protein-solvent system is generally described by a sum of the direct Coulombic energy and the dielectric shielding from the solvent due to the reaction field (42),

$$G_{\text{el}} = \frac{1}{2} \sum_{i \neq j} \frac{q_i q_j}{\epsilon_p |r_i - r_j|} + \frac{1}{2} \sum_i q_i \varphi^{\text{react}}(r_i) \quad (\text{Eq. 1})$$

The potential energy $\varphi^{\text{react}}(r_i)$ due to the dielectric shielding at the i th charge position r_i , can be given as the difference between the electrostatic potentials $\varphi^{\text{sol}}(r_i)$ and $\varphi^{\text{vac}}(r_i)$, both of which were calculated by numerically solving the Poisson equations,

$$\Delta \epsilon(r) \Delta \varphi(r) = -4\pi \sum_i q_i \delta(r - r_i) \quad (\text{Eq. 2})$$

assuming $\epsilon_p = 10$ and $\epsilon_p = \epsilon_s = 80$, respectively. Here, q_i is the i th charge and ϵ_p and ϵ_s are the dielectric constants $\epsilon(r)$ in the protein region and in the solvent region, respectively. ϵ_s in the solvent region was always assumed to be 80. The divergence of the electrostatic potential due to the self-Coulombic energy can be completely avoided, because the delta function in the right-hand side in eq. 2 is replaced by the smooth charge density in the actual numerical calculations (42).

In the continuum model for the native state, only the ionic charges were assumed as q_i to be located on both carboxyl oxygen atoms of the side chains of Asp and Glu, on the N ζ atoms of Lys, on the C ζ atoms of Arg, and on an N atom of the N terminus, respectively. When a strong ion pair is formed within 3 Å between the two heavy atoms, $-e$ charge was assumed on one of the two carboxyl oxygen atoms of Asp or Glu, which is the closest to the positively charged residue, and $+e$ charge was on N η 1, N η 2, or N ϵ , which is the closest to the negatively charged residue, in the case of Arg. Then, the above Poisson equations were numerically solved twice for the different ϵ_p values.

For the denatured state, the Coulombic term was calculated by using the root mean square distance $\langle r_{ij}^2 \rangle^{1/2}$ between the i th and j th charges in the different residues as the distance between those charges. We evaluated the mean square distance between the corresponding C α atoms for a simplified protein chain model, which was approximated by a Gaussian chain composed of only the connected C α atoms, with the mean inter-C α bond length 3.8 Å and the angle between the successive

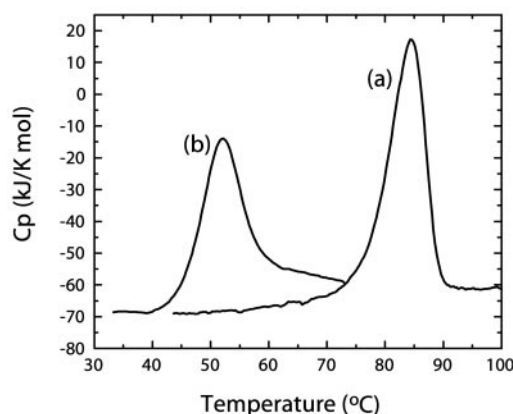


FIG. 1. Typical excess heat capacity curves of tryptophan synthase α -subunits from *P. furiosus* and *E. coli* near pH 9.5 in 20 mM Gly-KOH buffer. (a), *Pf*- α -subunit, at pH 9.60 and protein concentration of 0.40 mg/ml; (b), *Ec*- α -subunit, at pH 9.54, and protein concentration of 0.93 mg/ml.

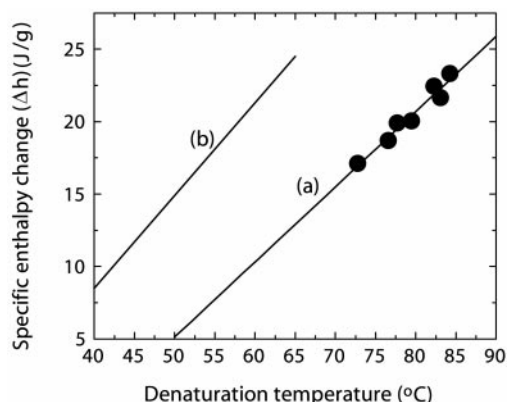


FIG. 2. Specific enthalpy change upon denaturation for tryptophan synthase α -subunit from the *P. furiosus* as a function of denaturation temperature. Filled circles represent each experimental point of *Pf*- α -subunit obtained at various pHs. Line (a) was given by a least-square fit to experimental points. Line (b) is a reported one for the *Ec*- α -subunit and the *St*- α -subunit (44).

three C α atoms 106.3° (43). Adding the squared distances between the C α atom and the charge position in each extended side chain for the *i*th and *j*th charges, respectively, r_{ij}^2 in the denatured state was estimated. As the value of $(\varphi^{\text{sol}}(r_i) - \varphi^{\text{vac}}(r_i))$ in the denatured state, the Poisson equations were solved for the models of the extended amino acid trimers, Ala-Xaa-Ala, where Xaa is Asp, Glu, Lys, or Arg. No charges were assumed on the N or C termini in the trimer models. The interaction term between the *i*th and *j*th charges in the reaction field calculation was neglected in the denatured states. A similar calculation was also performed for the N terminus using the extended Met-Ala model, assuming +e charge on the N atom of Met.

RESULTS

Thermal Stability of the α -Subunit of Tryptophan Synthase from *P. furiosus*—To examine the thermal stability of the *Pf*- α -subunit, DSC measurements were carried out at various pH values. Fig. 1 shows excess heat capacity curves of both the *Pf*- α -subunit and α -subunits from *E. coli* (*Ec*- α -subunit) near pH 9.5. The denaturation temperature of the *Pf*- α -subunit was about 33 °C higher than that of the *Ec*- α -subunit at the same conditions, indicating that the protein from a hyperthermophile has extremely high thermostability. In the acidic region from pH 3 to 4, the heat denaturation of the *Pf*- α -subunit was almost completely reversible, but the *Ec*- α -subunit was acid-denatured at room temperature. In Fig. 2 the specific enthalpy changes upon denaturation (Δh , the enthalpy value per a gram of proteins) obtained from DSC curves of the *Pf*- α -subunit at

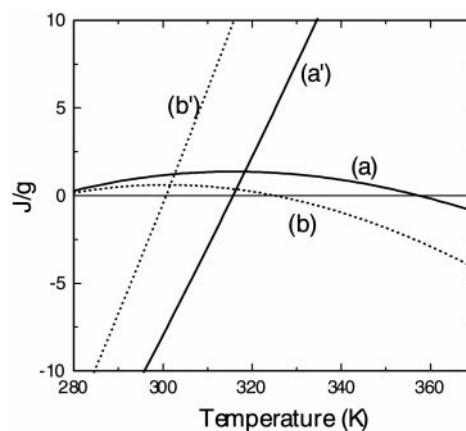


FIG. 3. Specific Gibbs energy change (ΔG) and specific entropy change ($T\Delta S$) upon denaturation of tryptophan synthase α -subunits from *P. furiosus* and *E. coli* as a function of temperature at pH 9.5. (a) and (a') represent ΔG and $T\Delta S$ of *Pf*- α -subunit, respectively. (b) and (b') represent ΔG and $T\Delta S$ of *Ec*- α -subunit, respectively.

various pH values are plotted against the denaturation temperature at each pH. Line (a) represents a least-square fit to the experimental points of the *Pf*- α -subunit. The slope corresponds to the specific heat capacity change of denaturation (Δc_p). The Δc_p value ($0.52 \text{ J K}^{-1} \text{ g}^{-1}$) of the *Pf*- α -subunit was slightly smaller than those of the *Ec*- α -subunit and the α -subunit from *S. typhimurium* (*St*- α -subunit), which are similar to each other ($0.64 \text{ J K}^{-1} \text{ g}^{-1}$ from the slope of line 2 in Sugisaki *et al.* (44)). The figure indicates that Δh values of the *Pf*- α -subunit were considerably lower than those of mesophile proteins at each temperature shown in the figure: for example, Δh was 10.3 J/g and 21.4 J/g for those from the *Pf*- α -subunit and mesophile proteins, respectively, at 60 °C. The thermodynamic parameters for heat denaturation as a function of temperature can be calculated using the following equations,

$$\Delta h(T) = \Delta h(T_d) - \Delta c_p(T_d - T) \quad (\text{Eq. 3})$$

$$\Delta s(T) = \Delta h(T_d)/T_d - \Delta c_p \ln(T_d/T) \quad (\text{Eq. 4})$$

$$\Delta g(T) = \Delta h(T) - T\Delta s(T) \quad (\text{Eq. 5})$$

assuming that Δc_p does not depend on temperature (45). The temperature functions of Δg for the *Pf*- α -subunit and the *Ec*- α -subunits at pH 9.5 obtained from eq. 5 are shown in Fig. 3, indicating that the *Pf*- α -subunit is more stable than the *Ec*- α -subunit under all the temperatures shown. The difference in Δg was remarkable at higher temperatures. Because the denaturation enthalpies of the *Pf*- α -subunit under all the temperatures as shown in Fig. 2 are lower than those of the *Ec*- α -subunit, the increase in Δg of *Pf*- α -subunit comes from the decrease in Δs as shown in Fig. 3.

Amino Acid Compositions and Overall Structure of the α -Subunit from *P. furiosus*—The *Pf*- α -subunit consists of 248 residues,² but the *St*- α -subunit has 268 (2). Table I shows the amino acid compositions of both α -subunits. Hydrophobic residues of the *Pf*- α -subunit were reduced from 157 (58.6%) to 133 (53.6%) as compared with those of the *St*- α -subunit: especially, Ala was 40 (14.9%) to 22 (8.9%). On the other hand, the hydrophilic residues were increased from 64 (23.9%) to 80 (32.3%); especially, Lys was 8 (3.0%) to 20 (8.1%). Jaenicke *et al.* (46) have reported that the ranking of the five most frequent amino acid exchanges from mesophiles to hyperthermophiles is seen to be Lys \rightarrow Arg, Ser \rightarrow Ala, Gly \rightarrow Ala, Ser \rightarrow Thr, and Ile \rightarrow Val. In the case of the *Pf*- α -subunit, however, Lys was remarkably increased and Ala decreased.

For the structure determination of the *Pf*- α -subunit, the

TABLE I
Comparison of amino acid compositions of tryptophan synthase α -subunits from *P. furiosus* and *S. typhimurium*

| Residues | α -Subunit from <i>P. furiosus</i> ^a | | α -Subunit from <i>S. typhimurium</i> ^a | |
|--------------|--|-------------------------------|---|-------------------------------|
| | Residue number | % of residue per total number | Residue number | % of residue per total number |
| Hydrophobic | 133 | 53.63 | 157 | 58.58 |
| Gly | 21 | 8.47 | 20 | 7.46 |
| Ala | 22 | 8.87 | 40 | 14.93 |
| Val | 21 | 8.47 | 17 | 6.34 |
| Leu | 25 | 10.08 | 28 | 10.45 |
| Ile | 17 | 6.85 | 18 | 6.72 |
| Met | 3 | 1.21 | 5 | 1.87 |
| Phe | 13 | 5.24 | 12 | 4.48 |
| Trp | 1 | 0.40 | 0 | 0.00 |
| Pro | 10 | 4.03 | 17 | 6.34 |
| Neutral | 35 | 14.11 | 47 | 17.54 |
| Ser | 11 | 4.44 | 16 | 5.97 |
| Thr | 14 | 5.65 | 7 | 2.61 |
| Asn | 7 | 2.82 | 11 | 4.10 |
| Gln | 2 | 0.81 | 10 | 3.73 |
| Cys | 1 | 0.40 | 3 | 1.12 |
| Hydrophilic | 80 | 32.26 | 64 | 23.88 |
| Asp | 12 | 4.84 | 13 | 4.85 |
| Glu | 20 | 8.06 | 16 | 5.97 |
| Lys | 20 | 8.06 | 8 | 2.99 |
| His | 4 | 1.61 | 5 | 1.87 |
| Arg | 15 | 6.05 | 15 | 5.60 |
| Tyr | 9 | 3.63 | 7 | 2.61 |
| Total number | 248 | | 268 | |

^a Amino acid compositions are taken from Ishida *et al.*² and Hyde *et al.* (2).

TABLE II
Data collection, phasing, and refinement statistics of tryptophan synthase α -subunit from *P. furiosus*

| Data set | Native | Hg | Pt |
|---|---------------|---------------|---------------|
| Data collection | | | |
| Wavelength (Å) | 1.0000 | 1.0000 | 1.5418 |
| Resolution range (Å) | 40.0–2.0 | 40.0–2.2 | 40.0–3.2 |
| Unique observations | 31,934 | 23,394 | 8,090 |
| Total observations | 90,544 | 83,106 | 24,473 |
| Completeness (%) | | | |
| Overall (final shell) | 94.2 (93.9) | 91.1 (93.9) | 95.9 (74.8) |
| R_{merge}^a | | | |
| Overall (final shell) | 0.035 (0.167) | 0.053 (0.216) | 0.092 (0.198) |
| MIRAS phasing | | | |
| R_{cullis}^b (acentric/centric) | | 0.75/0.81 | 0.92/0.91 |
| R_{cullis}^b —anomalous | | 0.77 | |
| Phasing power (acentric/centric) ^c | | 1.73/1.11 | 1.06/1.06 |
| Phasing power—anomalous | | 1.62 | |
| Figure of merit | | | 0.31 |
| Refinement statistics | | | |
| Resolution range (Å) | 40.0–2.0 | | |
| R -factor (%) ^d | 19.8 | | |
| R_{free} (%) | 24.9 | | |
| Root mean square deviations from ideality | | | |
| Bond distances (Å) | 0.006 | | |
| Bond angles (°) | 1.2 | | |

^a $R_{\text{merge}} = 100 \times \sum |I_{h,i} - \langle I_h \rangle| / \sum I_{h,i}$. $I_{h,i}$ are individual values, and $\langle I_h \rangle$ is the mean value of the intensity of reflection h .

^b $R_{\text{cullis}} = \langle \text{phase-integrated lack of closure} \rangle / \langle |F_{\text{ph}} - F_{\text{p}}| \rangle$.

^c Phasing power = $\langle |F_{\text{h}}(\text{cal})| / \text{phase-integrated lack of closure} \rangle$.

^d R -factor = $\sum |F_o| - |F_c| / \sum |F_o|$.

molecular replacement method using the structure of the α -subunit in the $\alpha_2\beta_2$ complex of tryptophan synthase from *S. typhimurium* (2) as a search model, was tried using AMoRe (47) or X-PLOR (39), because the sequence identity of both proteins was 31.5%. But all attempts to refine the solution failed. Therefore, the structure of the *Pf*- α -subunit was solved by the method of multiple isomorphous replacement using two derivatives with K_2PtCl_4 and CH_3HgCl (Table II). It was found that the *Pf*- α -subunit adopts an 8-fold α/β barrel fold, first observed in a triose-phosphate isomerase barrel (48), similar to that of the *St*- α -subunit (2), as shown in Figs. 4 and 6. The helices and

strands corresponding to canonical α/β barrel elements have been numbered consecutively 1 to 8. In the structure of the *St*- α -subunit, an extra helical segment, designated helix 0, precedes the first strand and acts as a cap in the bottom of the barrel (2). However, in the *Pf*- α -subunit the helix 0 was not observed because of the missing 12 residues in the N-terminal region as compared with the *St*- α -subunit. Six residues in the C-terminal helix 8 were also deleted in the *Pf*- α -subunit. Fig. 5 shows the secondary structure-based sequence alignment using the secondary structure elements assigned using DSSP (49). This alignment also suggests that another two residues

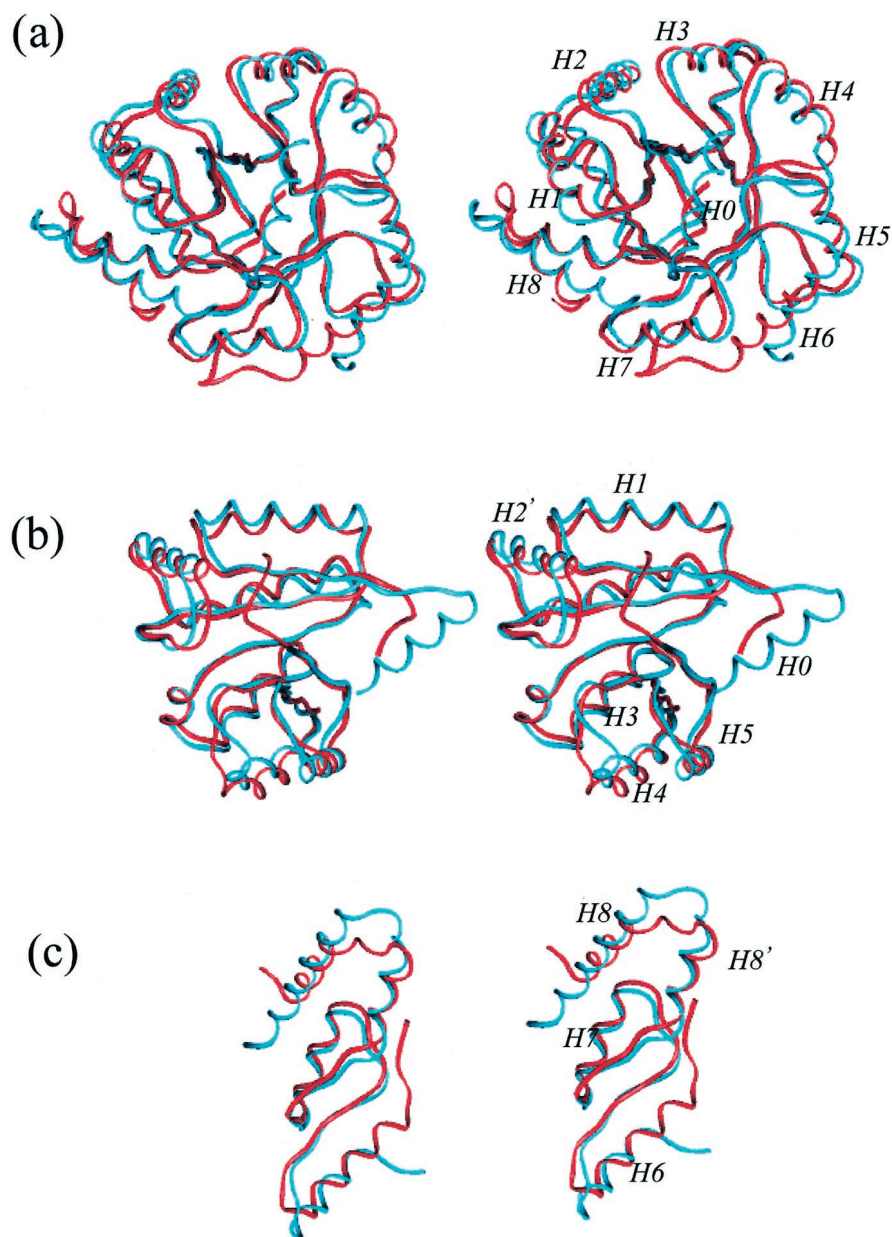


FIG. 6. Schematic stereo view of the α -subunits of tryptophan synthase from *P. furiosus* and from *S. typhimurium* superimposed. Red and blue represent the Pf- α -subunit and the St- α -subunit (1BKS), respectively. H0 through H8 represents helices 0 through 8 in the St- α -subunit, respectively. (a) represents the whole structures of both α -subunits. (b) and (c) represent stereo drawings of the main chains in the N- and C-terminal regions of tryptophan synthase α -subunits superimposed, respectively.

mational changes to less mobile forms. On the other hand the structure of the corresponding loop region of the Pf- α -subunit was remarkably different from that of the St- α -subunit with inhibitors (Fig. 6). This suggests that the loop region of the Pf- α -subunit might drastically change the conformation when the substrate (or inhibitor) binds to the protein.

Peaks II and IV in Fig. 7 are close to each residue corresponding to positions 41 and 95, respectively, in the St- α -subunit, which were deleted in the Pf- α -subunit. Peak III corresponds to helix 2'. In the case of the St- α -subunit a prominent deviation from the canonical triose-phosphate isomerase barrel is an insertion of 26 residues (positions 53–78) between strand 2 and helix 2. These residues, including helix 2', are highly conserved in α -subunits among many species. Two catalytic residues (Glu⁴⁹ and Asp⁶⁰) in the St- α -subunit are involved in or near the insertion region. The corresponding residues, Glu³⁶ and Asp⁴⁷, in the Pf- α -subunit were also found at coordinates similar to those from the mesophile protein. Helix 2' might be important in connection with enzyme function, although catalytic residues are not involved. In the region of peak V in the St- α -subunit, there are hydrogen-bonding residues (Val¹³³,

Glu¹³⁴, and Glu¹³⁵) interacting with the β -subunit (Fig. 5), which are important residues in the complex formation (53). The deviations of the peaks I, II, IV, and VII were caused by the deletion of residues, but other peaks of deviations (III, V, and VI) were related to enzymatic function, suggesting that the conformation of these portions might be affected when the α -subunit forms the complex with the β_2 -subunit.

High resolution structures of proteins from hyperthermophiles have shown that the number of ion pairs in most of the hyperthermophile proteins is higher than in mesophile counterparts (4–13), although additional ion pairs were not observed in some hyperthermophile proteins (25–28). Table III lists the ion pairs formed within 4 Å for the Pf- α -subunit and St- α -subunit. The number (0.05) of ion pairs per a residue for the Pf- α -subunit is relatively higher than that (0.02) for the St- α -subunit but is comparable to that (0.04) of the average number of mesophile proteins (54). In the Pf- α -subunit, 24 residues are involved in the formation of 13 ion pairs: five ion pairs for nine residues in the St- α -subunit. Only one ion pair, Arg⁵⁷-Asp¹⁵, in the Pf- α -subunit is conserved in the St- α -subunit (Arg⁷⁰-Asp²⁷). Ion pairs in the Pf- α -subunit form between

the following structural segments: five intra-helical, three inter-helical, two between helix and loop, one between helix and β -strand, one between β -strands, and one between β -strand and loop (Table III).

The volume of the cavities in the subunits of glutamate dehydrogenase from three different species decreases significantly with increasing thermal stability of the enzymes (9). For two α -subunits of tryptophan synthase, the cavity volume was determined by attempting to insert a probe sphere of 1.4-Å radius (assuming a water molecule) (55). In the case of the *Pf*- α -subunit, seven cavities were found and the total volume was 226.8 Å³; in the *St*- α -subunit eight cavities were found with 318.1 Å³ volume (Table IV). The energy term of ΔG due to changes in the cavity size can be expressed in terms of the cavity volume (100–140 J mol⁻¹ Å⁻³) (56). Using this parameter (120 J mol⁻¹ Å⁻³) the increment in stabilization of the *Pf*- α -subunit due to the decrease in cavity volume could be calculated to be 11.0 kJ/mol (Table IV), as compared with that of the *St*- α -subunit.

Because the *B*-factor in x-ray analysis can correlate with the flexibility of atoms, *B*-factors averaged for the main-chain atoms of the *Pf*- α -subunit were examined along with those of the *St*- α -subunit. Both data were collected at room temperatures (293–300 K). The difference in *B*-factors between both proteins (Fig. 8) shows that two parts of the *Pf*- α -subunit, corresponding to peaks I and II in Fig. 8, are more mobile than those of the *St*- α -subunit. The residues of peaks I and II are Tyr⁹³ and Phe¹²⁰, respectively. The corresponding residues in the *St*- α -subunit are Phe¹⁰⁷ and Glu¹³⁴, which are the residues (or neighbor residues) hydrogen bonding with the β_2 -subunit in the $\alpha_2\beta_2$ complex. This suggests that these residues (Tyr⁹³ and Phe¹²⁰) in the *Pf*- α -subunit might become less flexible when the

Pf- α -subunit associates with the β_2 -subunit. The *B*-factor values in other parts of the *Pf*- α -subunit were relatively lower than those of the *St*- α -subunit. This may be due to enhanced thermal stability of proteins from hyperthermophiles in the standard state at 25 °C, caused by enhanced conformational rigidity in their folded native state (30).

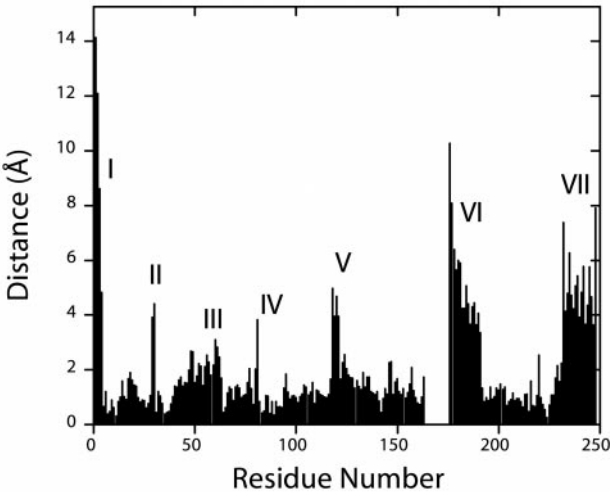


FIG. 7. Root mean square deviations (Å) of C α atoms between the *Pf*- α -subunit and the *St*- α -subunit after a least squares fit of corresponding C α atoms. Peaks I through VII represent only a discrimination peak of large differences. Residual numbers are for the *Pf*- α -subunit.

TABLE III
Ion pairs (salt bridges) in tryptophan synthase α -subunits from *P. furiosus* and *S. typhimurium* within 4 Å

| No. | Positively charged side chain | Accessibility | Negatively charged side chain | Accessibility | Location | Distance |
|-----------------------------------|-------------------------------|---------------|-------------------------------|---------------|----------|----------|
| | | % | | % | | Å |
| (A) <i>Pf</i> - α -subunit | | | | | | |
| 1 | Arg-57:N ϵ | 0 | Asp-15:O δ 2 | 2.4 | H2-S1 | 2.93 |
| | Arg-57:NH2 | 0.3 | Asp-15:O δ 1 | 0.1 | H2-S1 | 2.96 |
| | Arg-57:NH2 | 0.3 | Asp-15:O δ 2 | 2.4 | H2-S1 | 3.72 |
| 2 | Arg-66:NH2 | 25.8 | Glu-103:O ϵ 2 | 18.0 | L-H3' | 3.02 |
| 3 | Arg-76:NH2 | 0.5 | Asp-110:O δ 1 | 0 | H2'-L | 2.60 |
| | Arg-76:NH2 | 0.5 | Asp-110:O δ 2 | 81.3 | H2'-L | 3.73 |
| 4 | Arg-77:NH2 | 59.8 | Glu-74:O ϵ 1 | 35.9 | H2'-H2' | 3.49 |
| 5 | Arg-98:N ϵ | 37.2 | Glu-132:O ϵ 2 | 35.8 | H3'-H4 | 2.86 |
| | Arg-98:NH2 | 19.3 | Glu-132:O ϵ 2 | 35.8 | H3'-H4 | 3.34 |
| 6 | Arg-98:NH1 | 55.9 | Glu-131:O ϵ 2 | 9.8 | H3'-H4 | 3.46 |
| | Arg-98:NH2 | 19.3 | Glu-131:O ϵ 2 | 9.8 | H3'-H4 | 2.54 |
| 7 | Arg-184:NH1 | 43.0 | Asp-181:O δ 2 | 40.7 | H6-H6 | 2.57 |
| | Arg-184:NH2 | 48.0 | Asp-181:O δ 2 | 40.7 | H6-H6 | 3.25 |
| 8 | Arg-185:NH1 | 10.5 | Asp-181:O δ 1 | 43.5 | H6-H6 | 2.75 |
| 9 | Arg-188:NH1 | 19.6 | Asp-146:O δ 2 | 55.4 | H6-H5 | 3.82 |
| | Arg-188:NH2 | 33.2 | Asp-146:O δ 1 | 2.8 | H6-H5 | 3.97 |
| | Arg-188:NH2 | 33.2 | Asp-146:O δ 2 | 55.4 | H6-H5 | 2.76 |
| 10 | Arg-232:NH1 | 56.0 | Asp-17:O δ 2 | 15.0 | L-S1 | 2.53 |
| | Arg-232:NH1 | 56.0 | Asp-17:O δ 1 | 0.2 | L-S1 | 3.68 |
| | Arg-232:NH2 | 68.6 | Asp-17:O δ 1 | 0.2 | L-S1 | 3.99 |
| 11 | Lys-193:N ζ | 25.6 | Asp-4:O δ 2 | 0.3 | S7-S1 | 2.65 |
| 12 | Lys-225:N ζ | 85.7 | Glu-229:O ϵ 2 | 27.2 | H8'-H8' | 3.90 |
| 13 | Lys-241:N ζ | 28.7 | Glu-244:O ϵ 1 | 81.3 | H8'-H8' | 3.79 |
| | Lys-241:N ζ | 28.7 | Glu-244:O ϵ 2 | 42.0 | H8'-H8' | 3.47 |
| (B) <i>St</i> - α -subunit | | | | | | |
| 1 | Arg-3:NH1 | 25.5 | Glu-2:O ϵ 2 | 67.7 | H0-H0 | 3.82 |
| 2 | Arg-14:NH2 | 20.2 | Glu-16:O ϵ 2 | 1.5 | L-H0 | 3.04 |
| 3 | Arg-70:NH2 | 0 | Asp-27:O δ 2 | 0.1 | H2'-L | 2.86 |
| | Arg-70:NH2 | 0 | Asp-27:O δ 1 | 0 | H2'-L | 3.18 |
| 4 | His-92:N ϵ 2 | 0 | Asp-38:O δ 1 | 1.4 | L-H1 | 2.66 |
| 5 | Lys-91:N ζ | 26.8 | Asp-38:O δ 1 | 1.4 | L-H1 | 3.98 |
| | Lys-91:N ζ | 26.8 | Asp-38:O δ 2 | 38.2 | L-H1 | 3.23 |

TABLE IV

Estimate of the difference in stability between α -subunits from *P. furiosus* and *S. typhimurium* on the basis of structural information

ΔG_{HP} , ΔG_{CAV} , ΔG_{ENT} ($= -T\Delta S$), and ΔG_{E1} represent ΔG values due to hydrophobic interaction, cavity volume, entropic effect, and electrostatic interaction, respectively.

| | <i>Pf</i> - α -subunit | <i>St</i> - α -subunit | $\Delta (= Pf - St)$ |
|----------------------------------|-------------------------------|-------------------------------|-----------------------|
| ASA value in N state | | | |
| C/S atoms | 5,600 Å ² | 6,097 Å ² | -497 Å ² |
| N/O atoms | 5,515 Å ² | 5,150 Å ² | 365 Å ² |
| ASA value in D state | | | |
| C/S atoms | 22,049 Å ² | 23,015 Å ² | -966 Å ² |
| N/O atoms | 12,468 Å ² | 12,364 Å ² | 104 Å ² |
| Difference of ASA values (D-N) | | | |
| C/S atoms | 16,449 Å ² | 16,918 Å ² | -469 Å ² |
| N/O atoms | 6,953 Å ² | 7,214 Å ² | -261 Å ² |
| ΔG_{HP} | 2,837.3 kJ/mol | 2,917.1 kJ/mol | -79.8 kJ/mol |
| Cavity Vol. (probe 1.4 Å) | 226.75 Å ³ | 318.07 Å ³ | -91.32 Å ³ |
| ΔG_{CAV} | -27.2 kJ/mol | -38.2 kJ/mol | 11.0 kJ/mol |
| Total number of residues | 248 | 268 | -20 |
| ΔG_{ENT} at 25 °C | | | 51.2 kJ/mol |
| ΔG_{ENT} at 60 °C | | | 57.2 kJ/mol |
| Electrostatic energy (1) | | | |
| ΔG_{E1} | 80.4 kJ/mol | 37.7 kJ/mol | 42.7 kJ/mol |
| Electrostatic energy (2) | | | |
| Native state | | | |
| Interaction energy | -748.9 kJ/mol | -373.0 kJ/mol | |
| Reaction-field energy | -1,221.8 | -985.8 | |
| Total energy | -1,970.7 | -1,358.8 | |
| Denatured state | | | |
| Interaction energy | -13.8 | 3.3 | |
| Reaction-field energy | -2,232.3 | -1,639.4 | |
| Total energy | -2,246.0 | -1,636.2 | |
| ΔE_{EL} | -275.3 | -277.3 | 2.0 kJ/mol |

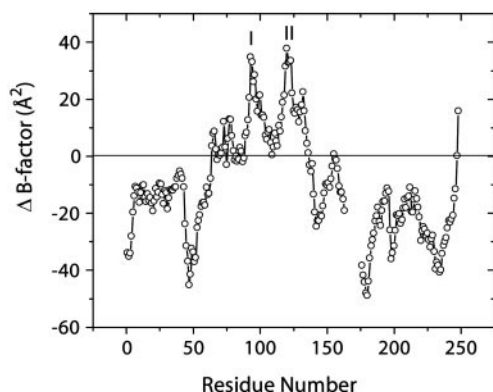


FIG. 8. Differences in B-factors (Å²) averaged for the main-chain atoms versus residue number between α -subunits of tryptophan synthases from *P. furiosus* and *S. typhimurium*. ΔB -factor subtracts the values of the *St*- α -subunit from those of the *Pf*- α -subunit. Peaks I and II correspond to Tyr⁹³ and Phe¹²⁰ of the *Pf*- α -subunit, respectively.

DISCUSSION

The α -subunit of tryptophan synthase from the hyperthermophile, *P. furiosus*, was remarkably stable as compared with those from mesophiles, *E. coli* and *S. typhimurium*. Calorimetric results indicated that the higher stability is caused by an entropic effect. This stabilization mechanism was investigated on the basis of structural differences between α -subunits of *P. furiosus* and *S. typhimurium*.

Contribution of the Hydrophobic Interaction—The hydrophobic interaction is one of the important stabilizing forces of folded protein structures (57). The hydrophobic effects have been extensively analyzed by studies with site-directed mutagenesis, and it has been revealed that hydrophobic residues in the interior of a protein contribute to the conformational stability (58–63). Takano *et al.* (64) have found a general rule

for the relationship between hydrophobic effect and conformational stability of a protein, using a series of hydrophobic mutants of human lysozyme. Furthermore, the change in unfolding Gibbs energy (ΔG) due to hydrophobic effect between wild-type and mutant proteins ($\Delta\Delta G_{\text{HP}}$) could be expressed using changes in accessible surface area (ASA) of nonpolar and polar atoms due to denaturation (65),

$$\Delta\Delta G_{\text{HP}} = \alpha\Delta\Delta\text{ASA}_{\text{nonpolar}} + \beta\Delta\Delta\text{ASA}_{\text{polar}} \quad (\text{Eq. 6})$$

where $\Delta\Delta\text{ASA}_{\text{nonpolar}}$ and $\Delta\Delta\text{ASA}_{\text{polar}}$ represent the differences in ΔASA of nonpolar and polar atoms of all residues in a protein, respectively, upon denaturation between wild-type and mutant proteins. Using the stability/structure data base for a series of mutant human lysozymes, the parameters, α and β , have been determined to be 0.178 and $-0.013 \text{ kJ mol}^{-1} \text{ \AA}^{-2}$, respectively (65). ASA values of proteins in the native state can be calculated using x-ray crystal structures, and those in the denatured state using their extended structures. For the calculation of ASA, C/S atoms in residues were assigned to $\text{ASA}_{\text{nonpolar}}$ and N/O to $\text{ASA}_{\text{polar}}$.

We tried to estimate the difference in contribution of the hydrophobic interaction between *Pf*- and *St*- α -subunits ($\Delta\Delta G_{\text{HP}}$). ASA values using x-ray crystal structures of the *Pf*- and *St*- α -subunits in the native state were calculated using the procedure of Connolly (55). The ASA values in the denatured states were calculated from their extended structures of both proteins, which were generated from the native structures using Insight II. As shown in Table IV, the ΔG values due to hydrophobic interaction (ΔG_{HP}) of the *Pf*- α -subunit was lower than that of *St*- α -subunit: the $\Delta\Delta G_{\text{HP}}$ value was -79.8 kJ/mol . This means that the higher stability of the *Pf*- α -subunit is not caused by the hydrophobic interaction. This fact could be speculated from the comparison of the contents of hydrophobic residues (Table I): the percent content of hydrophobic residues for the *Pf*- α -subunit (53.6%) was lower than that for the *St*- α -

subunit (58.65%). Makhatadze and Privalov (66) have reported that protein folding is an enthalpically driven process caused by van der Waals interactions between nonpolar groups in the interior of a protein. In the case of the *Pf*- α -subunit, the higher stability was not caused by enthalpic effects, suggesting that the hydrophobic interactions due to nonpolar groups in the interior of the protein do not dominate as compared with those of the *St*- α -subunit. This is also confirmed by the result that the ΔC_p value of the *Pf*- α -subunit ($0.52 \text{ J K}^{-1} \text{ g}^{-1}$) was lower than that for the *St*- α -subunit ($0.64 \text{ J K}^{-1} \text{ g}^{-1}$).

The ratio of hydrophobic residues (Val, Leu, and Ile) is generally higher in hyperthermophile proteins than in mesophile ones (67). But it is not certain that hydrophobic interactions are strengthened at higher temperatures. Thermodynamic analysis of hydrophobic interactions (66) indicates that at high temperature the entropy of hydration of a nonpolar group decreases to zero and the Gibbs energy of the hydrophobic interaction becomes completely enthalpic. This results in the Gibbs energy of the hydrophobic interactions reaching its maximum strength around 75°C , which is lower than the optimum growth temperatures of hyperthermophiles. This suggests that the hydrophobic interaction does not always act efficiently for the stabilization of a protein at extremely high temperatures over 100°C .

Contribution of Ion Pairs (Salt Bridge)—It seems that ion pairs (salt bridges) have important roles in protein stability, because they occur frequently in hyperthermophile proteins. However, Xiao and Honig (23) have reported that the roles of individual ion pairs and ion pair networks are variable, and among them some enhance protein stability, whereas others reduce it. The contribution depends strongly on the detailed environment of a particular network. Recently, using computation of ionic interaction energies by solving the Poisson equation in a continuum solvent medium, the ion pairs of glutamate dehydrogenase from *P. furiosus* have been reported to be highly important whereas those in the mesophilic homologs are only marginally stabilizing (13). These reports suggest the necessity to evaluate the strength of each ion pair in a protein. Therefore, we calculated electrostatic free energies in the *Pf*- α -subunit and the *St*- α -subunit solving the Poisson equations numerically for their continuum.

As shown in Table IV, the electrostatic free energy in the native state contributed by the direct interactions among ionic charges was much less in the *Pf*- α -subunit than that in the *St*- α -subunit. Because the *Pf*- α -subunit has more ionizable residues than the *St*- α -subunit, the reaction field energy in the native state of the *Pf*- α -subunit was also much less than that of the *St*- α -subunit. Thus, the ion pairs of the *Pf*- α -subunit greatly contribute to the conformational stability in the native state, as compared with those of the *St*- α -subunit.

However, the stability of the protein should be accounted for by the difference between the native state and the denatured state. Suppose that almost all the protein atoms in the denatured states are well exposed to the solvent molecules with the extended peptide chain conformations. The reaction field energy should become a very large negative value as the number of the ionizable residues increases. Namely, the *Pf*- α -subunit in the denatured state was also more stable than the *St*- α -subunit in the denatured state due to the reaction field energy. Consequently, as shown in the right-hand column of Table IV, $\Delta\Delta E$ between ΔE in the *Pf*- α -subunit and that in the *St*- α -subunit is only 2 kJ/mol , which may not explain all of the causes for the difference in the conformational stabilities of the two proteins.

In this continuum model, both absolute free energy values in the native states and in the denatured states were so large that even a slight deviation in the parameters associated with the

model would give a large error without revising the parameter values. In the current computation, it is not the comparison between the wild-type protein and the point mutants as once calculated by Kumar *et al.* (13) who neglected the electrostatic free energies in the denatured states. But it is the comparison between proteins from different species with very many substitutions of amino acid residues, and so the electrostatics in the denatured state cannot be neglected.

Recently, Takano *et al.* (24) have estimated the strength of an ionic interaction using a series of ion pair mutants of human lysozyme. Each contribution is not equivalent, but they have found that the contribution correlates linearly with the solvent-inaccessibility of the ion pairs; the contribution of ion pairs is small when 100% accessible, whereas it is about 9 kJ/mol if 100% inaccessible. Then, using accessibility of ion pair atoms in Table III, the contribution due to electrostatic interactions of the *Pf*- α -subunit and the *St*- α -subunit was calculated to be 80.4 and 37.7 kJ mol^{-1} , respectively (ΔG_{EL} in Table IV). This suggests that ion pairs in the *Pf*- α -subunit play an important role in conformational stability.

Contribution of the Entropic Effect—One way of increasing the stability of a protein ($\Delta G = \Delta H - T\Delta S$) is to lower its denaturation entropy (ΔS). The substitutions of other residues by less flexible residues, Pro or Gly (68–70), and the introduction of a disulfide bond (71) are confirmed to increase the stability of proteins. The conformational entropy of a protein in the denatured state can be also lowered by shortening the polypeptide chain. For thermophile proteins, a shortening of the N and C termini and the reduction of loop sizes have been observed (10, 28, 72, 73).

In the case of the *Pf*- α -subunit, 12 residues in the N-terminal, six in the C-terminal, and one residue each in two loops are deleted compared with the sequence of the *St*- α -subunit (Fig. 5). Oobatake and Ooi (74) have estimated the difference in thermodynamic parameters of each amino acid residue upon denaturation. Using these parameters the difference in contribution due to conformational entropy upon denaturation ($-T\Delta S$) between both proteins was calculated to be 51.2 kJ mol^{-1} and 57.2 kJ mol^{-1} , at 25 and 60°C , respectively (ΔG_{ENT} in Table IV). A great negative value of difference in enthalpy between both proteins ($\Delta\Delta H = -330.2 \text{ kJ mol}^{-1}$ at 60°C from Fig. 2) could be partly compensated for by the difference in entropic effect (57.2 kJ mol^{-1}). This indicates that the entropic effect due to these deletions plays an important role in the stabilization of the *Pf*- α -subunit. Recently, by analyzing many homologous relationships among many translated open reading frames of DNA from the 20 proteomes, Thompson and Eisenberg (75) have reported that there is a general trend in nature for thermophile sequences to be shorter than their mesophile homologs. This finding has not yet received as much attention as have ion pair networks (75). Present experimental results strongly support this transproteomic evidence.

The N-terminal segments of most prokaryotic tryptophan synthase α -subunits are approximately the same length and contain a helix 0. However, α -subunits from thermophiles, *Bacillus stearothermophilus* and *Thermotoga maritima*, appear to lack eight and 16 residues, respectively, at their N termini, similar to deletion of 12 residues in the *Pf*- α -subunit. These deletions might be related to the higher stability due to entropic effect.

Conclusion—The *Pf*- α -subunit from hyperthermophile, *P. furiosus*, was especially stable in the high temperature region as compared with homologous counterparts from mesophiles. DSC experiments indicated that its higher stability was caused by entropic factors. The tertiary structure of the *Pf*- α -subunit could be analyzed to interpret these thermodynamic character-

istics showing thermostabilization of the hyperthermophile protein. On the basis of structural information of the *Pf*- α -subunit and the *St*- α -subunit, it could be concluded that 1) the contribution of hydrophobic interaction of *Pf*- α -subunit to stability was considerably lower than that of the *St*- α -subunit, and 2) the higher stability of the *Pf*- α -subunit was caused by a decrease in cavity volume in the interior of the molecule, increase in ionic interaction (salt bridge), and entropic effects due to shortening of the polypeptide chain.

Acknowledgments—We greatly thank Prof. Tomitake Tsukihara (Institute for Protein Research, Osaka University) for valuable discussion on the x-ray analysis of the α -subunit structure of tryptophan synthase from *P. furiosus*.

REFERENCES

- Miles, E. W. (1995) *Subcell. Biochem.* **24**, pp. 207–254
- Hyde, C. C., Ahmed, S. A., Paduano, E. A., Miles, E. W., and Davies, D. R. (1988) *J. Biol. Chem.* **263**, 17857–17871
- Schultz, G. E., and Creighton, T. E. (1969) *Euro. J. Biochem.* **10**, 195–197
- Day, M. W., Hsu, B. T., Joshua-Tor, L., Park, J.-B., Zhou, Z. H., Adams, M. W. W., and Rees, D. C. (1992) *Protein Sci.* **1**, 1494–1507
- Chan, M. K., Mukund, S., Kletzin, A., Adams, M. W. W., and Rees, D. C. (1995) *Science* **267**, 1463–1469
- Hennig, M., Darimont, B., Sterner, R., Kirschner, K., and Jansonius, J. N. (1995) *Structure* **3**, 1295–1306
- Yip, K. S., Stillman, T. J., Britton, K. L., Artymiuk, P. J., Baker, P. J., Sedelnikova, S. E., Engel, P. C., Pasquo, A., Chiaraluce, R., Conslvi, V., Scandurra, R., and Rice, D. W. (1995) *Structure* **2**, 1147–1158
- Dedecker, B. S., O'Brien, R., Fleming, P. J., Geider, J. H., Jackson, S. P., and Sigler, P. B. (1996) *J. Mol. Biol.* **264**, 1072–1084
- Knapp, S., de Vos, W. M., Rice, D., and Ladenstein, R. (1997) *J. Mol. Biol.* **267**, 916–932
- Tahirov, H., Oki, H., Tsukihara, T., Ogasahara, K., Yutani, K., Ogata, K., Izu, Y., Tsunasawa, S., and Kato, I. (1998) *J. Mol. Biol.* **284**, 101–124
- Russell, R. J. M., Ferguson, J. M. C., Hough, D. W., Danson, M. J., and Taylor, G. L. (1997) *Biochemistry* **36**, 9983–9994
- Isupov, M. N., Fleming, T. M., Dalby, A. R., Crowhurst, G. S., Bourne, P. C., and Littlechild, J. A. (1999) *J. Mol. Biol.* **291**, 651–660
- Kumar, S., Ma, B., Tsai, C.-J., and Nussionv, R. (2000) *Proteins* **38**, 368–383
- Horovitz, A., Serrano, L., Avron, B., Bycroft, M., and Fersht, A. R. (1990) *J. Mol. Biol.* **216**, 1031–1044
- Serrano, L., Horovitz, A., Avron, B., Bycroft, M., and Fersht, A. R. (1990) *Biochemistry* **29**, 9343–9352
- Erwin, C. R., Barnett, B. L., Oliver, J. D., and Sullivan, J. F. (1990) *Protein Eng.* **4**, 87–97
- Sali, D., Bycroft, M., and Fersht, A. R. (1991) *J. Mol. Biol.* **220**, 779–788
- Dao-pin, S., Sauer, U., Nicholson, H., and Matthews, B. W. (1991) *Biochemistry* **30**, 7142–7153
- Lyu, P. C., Gans, P. J., and Kallenbach, N. R. (1992) *J. Mol. Biol.* **223**, 343–350
- Vetriani, C., Maeder, D. L., Toliaday, N., Yip, K. S., Stillman, T. J., Britton, K. L., Rice, D. W., Klump, H. H., and Robb, F. (1998) *Proc. Natl. Acad. Sci. U. S. A.* **95**, 12300–12305
- Spek, E. J., Bui, A. H., Lu, M., and Kallenbach, N. R. (1998) *Protein Sci.* **7**, 2431–2437
- Sanz-Aparicio, J., Hermoso, J. A., Martinez-Ripoll, M., Gonzalez, B., Lopez-Camacho, C., and Polaina, J. (1998) *Proteins Struct. Funct. Genet.* **33**, 567–576
- Xiao, L., and Honig, B. (1999) *J. Mol. Biol.* **289**, 1435–1444
- Takano, K., Tsuchimori, K., Yamagata, Y., and Yutani, K. (2000) *Biochemistry* **39**, 12375–12381
- Macedo-Riberio, S., Darimont, B., Sterner, R., and Huber, R. (1996) *Structure* **4**, 1291–1301
- Edmonson, S. P., Qui, L., and Shriver, J. W. (1995) *Biochemistry* **34**, 13289–13304
- Cavagnero, S., Zhou, Z. H., Adams, M. W., and Chan, S. I. (1995) *Biochemistry* **34**, 9865–9873
- Starich, M. R., Sandman, K., Reeve, J. N., and Summers, M. (1996) *J. Mol. Biol.* **255**, 187–203
- Pace, C. N., Shirley, B. A., McNutt, M., and Gajiwala, K. (1996) *FASEB. J.* **10**, 75–83
- Jaenicke, R., and Bohm, G. (1998) *Curr. Opin. Struct. Biol.* **8**, 738–748
- Merritt, E. A., and Bacon, D. J. (1997) *Methods Enzymol.* **277**, 505–524
- Sakabe, N. (1991) *Nucl. Instr. Methods Phys. Res.* **A303**, 448–463
- Otwinowski, Z., and Minor, W. (1997) *Methods Enzymol.* **276**, 307–326
- Matthews, B. W. (1968) *J. Mol. Biol.* **33**, 491–497
- Collaborative Computational Project, Number 4. (1994) *Acta Crystallogr. Sect. D Biol. Crystallogr.* **50**, 760–763
- de La Fortelle, E., and Bricogne, G. (1997) *Methods Enzymol.* **276**, 472–493
- Abrahams, J. P. (1996) *Acta Crystallogr. Sect. D Biol. Crystallogr.* **52**, 30–42
- Jones, T. A., Zou, J. Y., Cowan, S. W., and Kjeldgaard, M. (1991) *Acta Crystallogr. Sect. A* **47**, 110–119
- Brunger, A. T. (1993) *X-plor Version 3.8.5.1* Yale University Press, New Haven, CT
- Brunger, A. T., Adams, P. D., Clore, G. M., DeLano, W. L., Gros, P., Grosse-Kunstleve, R. W., Jiang, J.-S., Kuszewski, J., Nilges, M., Pannu, N. S., Read, R. J., Rice, L. M., Simonson, T., & Warren, G. L. (1998) *Acta Cryst. D54*, 905–921
- Laskowski, R. A., McArthur, M. W., Moss, D. S., and Thornton, J. M. (1993) *J. Appl. Cryst.* **26**, 283–291
- Nakamura, H. (1996) *Rev. Biophys.* **29**, 1–90
- Levitt, M. (1976) *J. Mol. Biol.* **104**, 59–107
- Sugisaki, Y., Ogasahara, K., Miles, E. W., and Yutani, K. (1990) *Thermo-chimica Acta* **163**, 117–122
- Privalov, P. L., and Khechinashvili, N. N. (1974) *J. Mol. Biol.* **86**, 665–684
- Jaenicke, R., Schurig, H., Beaucamp, N., and Ostendorp, R. (1996) *Adv. Protein Chem.* **48**, 181–270
- Navaza, J. (1994) *Acta Crystallogr. Sect. D Biol. Crystallogr.* **50**, 157–163
- Banner, D. W., Bloomer, A. C., Petsko, G. A., Phillips, D. C., and Wilson, I. A. (1976) *Biochem. Biophys. Res. Commun.* **72**, 146–155
- Kabsch, W., and Sander, C. (1983) *Biopolymers* **22**, 2577–2637
- Rhee, S. Parris, K. D., Hyde, C. C., Ahmed, S. A., Miles, E. W., and Davies, D. R. (1997) *Biochemistry* **36**, 7664–7680
- Weyand, M., and Schlichting, I. (1999) *Biochemistry* **38**, 16469–16480
- Schneider, T. R., Gerhardt, E., Lee, M., Liang, P. H., Anderson, K. S., and Schlichting, I. (1998) *Biochemistry* **37**, 5394–5406
- Hiraga, K., and Yutani, K. (1997) *J. Biol. Chem.* **272**, 4935–4940
- Barlow, D. J., and Thornton, J. M. (1983) *J. Mol. Biol.* **168**, 867–885
- Connolly, M. L. (1993) *J. Mol. Graphics* **11**, 139–141
- Eriksson, A. E., Baase, W. A., Zhang, X.-J., Heinz, D. W., Blaber, M., Baldwin, E. P., and Matthews, B. W. (1992) *Science* **225**, 178–183
- Kauzman, W. (1959) *Adv. Protein Chem.* **14**, 1–63
- Yutani, K., Ogasahara, K., Sugino, Y., and Matsusiro, A. (1977) *Nature* **267**, 274–275
- Yutani, K., Ogasahara, K., Tsujita, T., and Sugino, Y. (1987) *Proc. Natl. Acad. Sci. U. S. A.* **84**, 4441–4444
- Kellis, J. T., Nyberg, K., Sali, D., and Fersht, A. R. (1988) *Nature* **333**, 784–786
- Matsumura, M., Becktel, W. J., and Matthews, B. W. (1988) *Nature* **334**, 406–410
- Shortle, D., Stites, W. E., and Meeker, A. K. (1990) *Biochemistry* **30**, 9686–9697
- Sandberg, W. S., and Terwilliger, T. C. (1991) *Proc. Natl. Acad. Sci. U. S. A.* **88**, 1706–1710
- Takano, K., Yamagata, Y., and Yutani, K. (1998) *J. Mol. Biol.* **280**, 749–761
- Funahashi, J., Takano, K., Yamagata, Y., and Yutani, K. (1999) *Protein Eng.* **12**, 841–850
- Makhatadze, G. I., and Privalov, P. L. (1995) *Adv. Protein. Chem.* **47**, 307–425
- Szilagy, A., and Zavodszky, P. (2000) *Structure* **8**, 493–504
- Hecht, M., Sturtevant, J., and Sauer, R. (1986) *Proteins* **1**, 43–46
- Herning, T., Yutani, K., Inada, K., Kuroki, R., Matsushima, M., and Kikuchi, M. (1992) *Biochemistry* **31**, 7077–7085
- Nicholson, H., Tronrud, D., Becktel, W., Matthews, B. (1992) *Biopolymers* **32**, 1421–1441
- Matsumura, M., Becktel, W. J., Levitt, M., and Matthews, B. W. (1989) *Proc. Natl. Acad. Sci. U. S. A.* **86**, 6562–6566
- Russell, R. J. M., Hough, D. W., Danson, M. J., and Taylor, G. L. (1994) *Structure* **2**, 1157–1167
- Usher, K. C., de la Cruz, A. F., Dahlquist, F. W., Swanson, R. V., Simon, M. I., Remington, S. J. (1998) *Protein Sci.* **7**, 403–412
- Oobatake, M., and Ooi, T. (1993) *Prog. Biophys. Mol. Biol.* **59**, 237–284
- Thompson, M. J., and Eisenberg, D. (1999) *J. Mol. Biol.* **290**, 594–604
- Kraulis, P. J. (1991) *J. Appl. Crystallogr.* **24**, 946–950

Entropic Stabilization of the Tryptophan Synthase α -Subunit from a Hyperthermophile, *Pyrococcus furiosus* : X-RAY ANALYSIS AND CALORIMETRY

Yuriko Yamagata, Kyoko Ogasahara, Yusaku Hioki, Soo Jae Lee, Atsushi Nakagawa, Haruki Nakamura, Masami Ishida, Seiki Kuramitsu and Katsuhide Yutani

J. Biol. Chem. 2001, 276:11062-11071.

doi: 10.1074/jbc.M009987200 originally published online December 15, 2000

Access the most updated version of this article at doi: [10.1074/jbc.M009987200](https://doi.org/10.1074/jbc.M009987200)

Alerts:

- [When this article is cited](#)
- [When a correction for this article is posted](#)

[Click here](#) to choose from all of JBC's e-mail alerts

This article cites 75 references, 7 of which can be accessed free at <http://www.jbc.org/content/276/14/11062.full.html#ref-list-1>